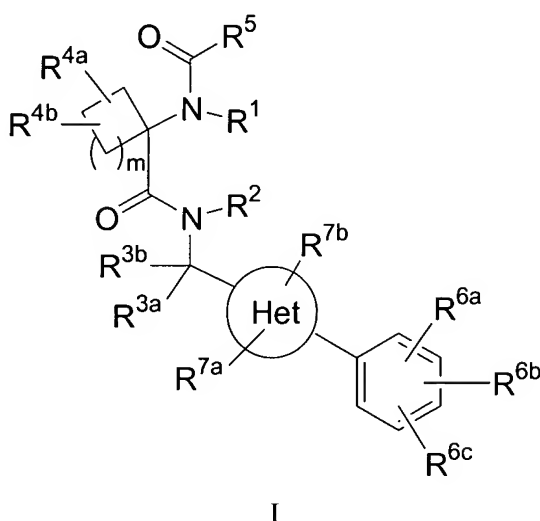


**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1 (original). A compound of formula I and pharmaceutically acceptable salts thereof:



wherein

Het is pyrimidinyl or pyridyl, or N-oxide thereof;

R<sup>1</sup> and R<sup>2</sup> are independently selected from hydrogen and C<sub>1-4</sub> alkyl;

R<sup>3a</sup> and R<sup>3b</sup> are independently selected from hydrogen and C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms;

R<sup>4a</sup> and R<sup>4b</sup> are independently selected from (1) hydrogen, (2) halogen, and (3) C<sub>1-4</sub> alkyl optionally substituted with 1 to 4 groups selected from halogen, OR<sup>a</sup>, OC(O)R<sup>a</sup>, S(O)<sub>k</sub>R<sup>d</sup>, OS(O)<sub>2</sub>R<sup>d</sup>, and NR<sup>1</sup>R<sup>2</sup>, or

R<sup>4a</sup> and R<sup>4b</sup> together with the carbon atom to which they are both attached form an exo-cyclic methylene optionally substituted with 1 to 2 groups selected from C<sub>1-4</sub> alkyl optionally substituted with 1-5 halogen atoms and C<sub>1-4</sub> alkyloxy;

R<sup>5</sup> is selected from (1) C<sub>1-6</sub> alkyl optionally substituted with 1 to 5 groups independently selected from halogen, nitro, cyano, OR<sup>a</sup>, SR<sup>a</sup>, COR<sup>a</sup>, SO<sub>2</sub>R<sup>d</sup>, CO<sub>2</sub>R<sup>a</sup>, OC(O)R<sup>a</sup>, NR<sup>b</sup>R<sup>c</sup>, NR<sup>b</sup>C(O)R<sup>a</sup>,

$\text{NR}^b\text{C}(\text{O})_2\text{R}^a$ ,  $\text{C}(\text{O})\text{NR}^b\text{R}^c$ ,  $\text{C}_{3-8}$  cycloalkyl, (2)  $\text{C}_{3-8}$  cycloalkyl optionally substituted with 1 to 5 groups independently selected from halogen, nitro, cyano and phenyl, (3)  $\text{C}_{3-6}$  alkynyl, (4)  $\text{C}_{2-6}$  alkenyl optionally substituted with hydroxyethyl, (5)  $(\text{CH}_2)_k$ -aryl optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano,  $\text{OR}^a$ ,  $\text{SR}^a$ ,  $\text{C}(\text{O})_2\text{R}^a$ ,  $\text{C}_{1-4}$  alkyl and  $\text{C}_{1-3}$  haloalkyl, wherein aryl is selected from phenyl, 3,4-methylenedioxyphenyl and naphthyl, (6)  $(\text{CH}_2)_k$ -heterocycle optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano,  $\text{OR}^a$ ,  $\text{SR}^a$ ,  $\text{C}_{1-4}$  alkyl and  $\text{C}_{1-3}$  haloalkyl wherein said heterocycle is selected from (a) a 5-membered heteroaromatic ring having a ring heteroatom selected from N, O and S, and optionally having up to 3 additional ring nitrogen atoms wherein said ring is optionally benzo-fused; (b) a 6-membered heteroaromatic ring containing from 1 to 3 ring nitrogen atoms and N-oxides thereof, wherein said ring is optionally benzo-fused; and (c) a 5- or 6-membered non-aromatic heterocyclic ring selected from tetrahydrofuranyl, 5-oxotetrahydrofuranyl, 2-oxo-2H-pyranyl, and 6-oxo-1,6-dihydropyridazinyl, (7)  $\text{C}(\text{O})_2\text{R}^a$ , (8)  $\text{C}(\text{O})\text{NR}^b\text{R}^c$ , and (9)  $\text{NR}^b\text{CO}_2\text{R}^a$ ;

$\text{R}^{6a}$  is selected from (1)  $\text{C}_{1-8}$  alkyl optionally substituted with 1-5 groups independently selected from halogen, nitro, cyano,  $\text{COR}^a$ ,  $\text{CO}_2\text{R}^a$ ,  $\text{C}(\text{O})\text{NR}^b\text{R}^c$ ,  $\text{OR}^a$ ,  $\text{OC}(\text{O})\text{R}^a$ ,  $\text{SR}^a$ ,  $\text{SO}_2\text{R}^d$ ,  $\text{S}(\text{O})\text{R}^d$ ,  $\text{NR}^b\text{R}^c$ ,  $\text{NR}^b\text{C}(\text{O})\text{R}^a$ ,  $\text{NR}^b\text{SO}_2\text{R}^d$ ,  $\text{NR}^b\text{CO}_2\text{R}^a$ , (2)  $\text{C}_{3-8}$  cycloalkyl, (3)  $\text{C}_{2-8}$  alkenyl optionally substituted with  $\text{CO}_2\text{R}^a$ , (4) halogen, (5) cyano, (6) nitro, (7)  $\text{NR}^b\text{R}^c$ , (8)  $\text{NR}^b\text{C}(\text{O})\text{R}^a$ , (9)  $\text{NR}^b\text{CO}_2\text{R}^a$ , (10)  $\text{NR}^b\text{C}(\text{O})\text{NR}^b\text{R}^c$ , (11)  $\text{NR}^b\text{C}(\text{O})\text{NR}^b\text{CO}_2\text{R}^a$ , (12)  $\text{NR}^b\text{SO}_2\text{R}^d$ , (13)  $\text{CO}_2\text{R}^a$ , (14)  $\text{COR}^a$ , (15)  $\text{C}(\text{O})\text{NR}^b\text{R}^c$ , (16)  $\text{C}(\text{O})\text{NHOR}^a$ , (17)  $\text{C}(=\text{NOR}^a)\text{R}^a$ , (18)  $\text{C}(=\text{NOR}^a)\text{NR}^b\text{R}^c$ , (19)  $\text{OR}^a$ , (20)  $\text{OC}(\text{O})_k\text{R}^a$ , (21)  $\text{S}(\text{O})_k\text{R}^d$ , (22)  $\text{SO}_2\text{NR}^b\text{R}^c$ , and (23) optionally substituted heterocycle where the heterocycle is selected from (a) a 5-membered heteroaromatic ring having a ring heteroatom selected from N, O and S, and optionally having up to 3 additional ring nitrogen atoms, (b) 4,5-dihydro-oxazolyl, and (3) 4,5-dihydro-1,2,4-oxadiazolyl, and wherein said substituent is 1 to 3 groups independently selected from  $\text{C}_{1-4}$  alkyl optionally substituted with 1 to 5 halogen atoms,  $\text{OR}^a$  or  $\text{OC}(\text{O})\text{R}^a$ ,

$\text{R}^{6b}$  and  $\text{R}^{6c}$  are independently selected from hydrogen, and a group from  $\text{R}^{6a}$ ; with the proviso that not more than one of  $\text{R}^{6a}$ ,  $\text{R}^{6b}$ , and  $\text{R}^{6c}$  is a heterocycle;

$\text{R}^{7a}$  and  $\text{R}^{7b}$  are independently selected from hydrogen, halogen, cyano, nitro,  $\text{OR}^a$ ,  $\text{CO}_2\text{R}^a$ ,  $\text{C}(\text{O})\text{NR}^b\text{R}^c$ ,  $\text{SO}_2\text{R}^d$ ,  $\text{NR}^b\text{R}^c$ , and  $\text{C}_{1-4}$  alkyl optionally substituted with 1 to 5 halogen atoms;

$\text{R}^a$  is selected from (1) hydrogen, (2)  $\text{C}_{1-4}$  alkyl optionally substituted with 1 to 5 halogen atoms, (3) phenyl optionally substituted with 1 to 3 groups independently selected from halogen, cyano, nitro,

OH, C<sub>1-4</sub> alkyloxy, C<sub>3-6</sub> cycloalkyl and C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, (4) C<sub>3-6</sub> cycloalkyl, and (5) pyridyl;

R<sup>b</sup> and R<sup>c</sup> are independently selected from (1) hydrogen, (2) C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 groups independently selected from halogen, amino, mono-C<sub>1-4</sub>alkylamino, di-C<sub>1-4</sub>alkylamino, and SO<sub>2</sub>R<sup>d</sup>, (3) (CH<sub>2</sub>)<sub>k</sub>-phenyl optionally substituted with 1 to 3 groups selected from halogen, cyano, nitro, OH, C<sub>1-4</sub> alkyloxy, C<sub>3-6</sub> cycloalkyl and C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, and (4) C<sub>3-6</sub> cycloalkyl, or

R<sup>b</sup> and R<sup>c</sup> together with the nitrogen atom to which they are attached form a 4-, 5-, or 6-membered ring optionally containing an additional heteroatom selected from N, O, and S; or

R<sup>b</sup> and R<sup>c</sup> together with the nitrogen atom to which they are attached form a cyclic imide;

R<sup>d</sup> is selected from (1) C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, (2) C<sub>1-4</sub> alkyloxy, and (3) phenyl optionally substituted with 1 to 3 groups selected from halogen, cyano, nitro, OH, C<sub>1-4</sub> alkyloxy, C<sub>3-6</sub> cycloalkyl and C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms;

k is 0, 1 or 2; and

m is 0, 1, 2 or 3.

2 (currently amended). A compound of Claim 1 wherein R<sup>5</sup> is (1) C<sub>1-6</sub> alkyl optionally substituted with 1 to 5 groups independently selected from halogen, nitro, cyano, OR<sup>a</sup>, SR<sup>a</sup>, COR<sup>a</sup>, SO<sub>2</sub>R<sup>d</sup>, CO<sub>2</sub>R<sup>a</sup>, OC(O)R<sup>a</sup>, NR<sup>b</sup>R<sup>c</sup>, NR<sup>b</sup>C(O)R<sup>a</sup>, C(O)NR<sup>b</sup>R<sup>c</sup>, and C<sub>3-8</sub> cycloalkyl, (2) 1,2,5-thiadiazolyl, (3) isoxazolyl, (4) isothiazolyl or (5) pyrimidinyl.

3 (currently amended). A compound of Claim 1 wherein R<sup>5</sup> is C<sub>1-3</sub> alkyl optionally substituted with 1 to 5 ~~group~~-halogen atoms wherein said halogen is chloro or fluoro.

4 (original). A compound of Claim 1 wherein R<sup>5</sup> is selected from difluoromethyl, dichloromethyl, chlorodifluoromethyl, trifluoromethyl, 1,1-dichloroethyl and 2,2,2-trifluoroethyl.

5 (original). A compound of Claim 1 wherein R<sup>5</sup> is pyrimidinyl.

6 (original). A compound of Claim 1 wherein R<sup>5</sup> is 1,2,5-thiadiazolyl, isoxazolyl or isothiazolyl.

7 (original). A compound of Claim 1 wherein R<sup>6a</sup> is OR<sup>a</sup>, CO<sub>2</sub>R<sup>a</sup> or tetrazolyl optionally substituted with C<sub>1-4</sub> alkyl.

8 (original). A compound of Claim 1 wherein R<sup>6a</sup> is OR<sup>a</sup>, CO<sub>2</sub>R<sup>a</sup> or tetrazolyl optionally substituted with C<sub>1-4</sub> alkyl, R<sup>6b</sup> is hydrogen or halogen, and R<sup>6c</sup> is hydrogen or halogen.

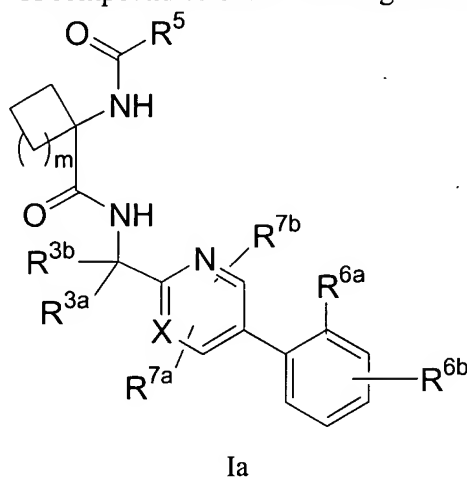
9 (original). A compound of Claim 1 wherein R<sup>6a</sup> is methoxycarbonyl, ethoxycarbonyl, C<sub>1-4</sub>alkoxy optionally substituted with 1 to 5 halogen atoms, or 2-methyl-2H-tetrazol-5-yl, R<sup>6b</sup> is fluoro or chloro, and R<sup>6c</sup> is hydrogen, chloro or fluoro.

10 (original). A compound of Claim 1 wherein Het is 2,5-pyridinediyl and R<sup>7a</sup> and R<sup>7b</sup> are independently hydrogen or halogen.

11 (original). A compound of Claim 10 wherein one of R<sup>7a</sup> and R<sup>7b</sup> is hydrogen and the other is fluoro or chloro.

12 (original). A compound of Claim 1 wherein m is 0 or 1.

13 (original). A compound of Claim 1 having formula Ia:



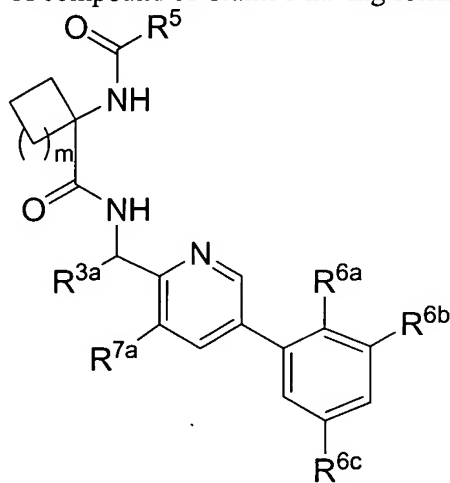
wherein X is carbon or nitrogen, and all other variables are as defined Claim 1.

14 (original). A compound of Claim 13 wherein m is 0 or 1 and one of R<sup>3a</sup> and R<sup>3b</sup> is hydrogen and the other is hydrogen or C<sub>1-3</sub>alkyl.

15 (original). A compound of Claim 13 wherein X is carbon, R<sup>7a</sup> is hydrogen or chloro or fluoro, and R<sup>7b</sup> is hydrogen.

16 (original). A compound of Claim 13 wherein X is nitrogen and R<sup>7a</sup> and R<sup>7b</sup> are each hydrogen.

17 (original). A compound of Claim 1 having formula Ib:

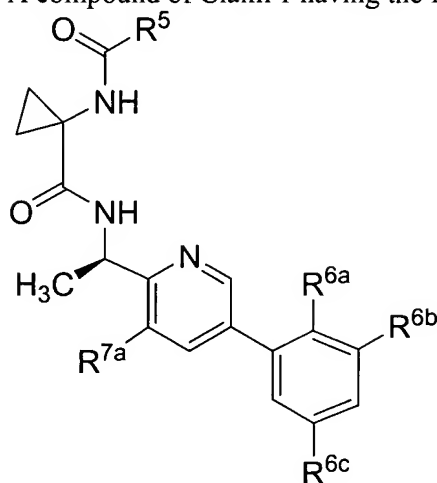


wherein m is 0 or 1, R<sup>3a</sup> is hydrogen or methyl, R<sup>6b</sup> and R<sup>6c</sup> are independently hydrogen, chloro or fluoro, R<sup>7a</sup> is hydrogen, chloro or fluoro, and the other variables are as defined in Claim 1.

18 (original). A compound of Claim 17 wherein R<sup>3a</sup> is hydrogen, and R<sup>6b</sup> and R<sup>7a</sup> are each independently chloro or fluoro.

19 (original). A compound of Claim 17 wherein R<sup>3a</sup> is hydrogen, R<sup>6b</sup> and R<sup>7a</sup> are each independently chloro or fluoro, R<sup>5</sup> is selected from isoxazolyl, thiazolyl, 1,2,5-thiadiazolyl, 5-pyrimidinyl and C<sub>1-2</sub>alkyl substituted with 1 to 3 halogen atoms selected from chloro and fluoro, and R<sup>6a</sup> is OR<sup>a</sup>, CO<sub>2</sub>R<sup>a</sup> or 2-methyl-5-tetrazolyl.

20 (original). A compound of Claim 1 having the formula Ic:



Ic

wherein R<sup>5</sup>, R<sup>6a</sup>, R<sup>6b</sup>, R<sup>6c</sup> and R<sup>7a</sup> are as defined in Claim 1.

21 (original). A compound of Claim 20 wherein R<sup>6b</sup> is halogen, and R<sup>6c</sup> and R<sup>7a</sup> are independently hydrogen or halogen.

22 (original). A compound of Claim 20 wherein R<sup>5</sup> is selected from isoxazolyl, isothiazolyl, 1,2,5-thiadiazolyl, 5-pyrimidinyl and C<sub>1-2</sub>alkyl substituted with 1 to 5 halogen atoms.

23 (original). A compound of Claim 21 wherein R<sup>5</sup> is selected from isoxazolyl, isothiazolyl, 1,2,5-thiadiazolyl, 5-pyrimidinyl and C<sub>1-2</sub>alkyl substituted with 1 to 5 halogen atoms.

24 (original). A compound of Claim 20 wherein R<sup>6a</sup> is selected from CO<sub>2</sub>C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy optionally substituted with 1 to 5 halogen atoms and 2-methyl-5-tetrazolyl.

25 (original). A compound of Claim 23 wherein R<sup>6a</sup> is selected from CO<sub>2</sub>C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy optionally substituted with 1 to 5 halogen atoms and 2-methyl-5-tetrazolyl.

26 (original). A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1 and pharmaceutically acceptable excipients.

27-32 (cancelled).